(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization

International Bureau



(43) International Publication Date 31 December 2003 (31.12.2003)

PCT

(10) International Publication Number WO 2004/000948 A1

(51) International Patent Classification7: 41/00, C09C 1/42

C09B 63/00,

(21) International Application Number:

PCT/IL2003/000497

(22) International Filing Date:

12 June 2003 (12.06.2003)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

150317

19 June 2002 (19.06.2002)

- (71) Applicant (for all designated States except US): YISSUM RESEARCH DEVELOPMENT COMPANY OF THE HEBREW UNIVERSITY OF JERUSALEM [IL/IL]; P.O. Box 39135 Givat Ram, 91390 Jerusalem (IL).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): YARIV, Shmuel [IL/IL]; 32 Maale Benjamin Street, 90620 Almon-Anatot D.N., East Binyamin (IL). LAPIDES, Isaak [IL/IL]; 1/16 Alley Qezia Gilo, Jerusalem 93754 (IL). PELED, Hagai [IL/IL]; 20 Alroi Street, Tel Aviv 65147 (IL). YIRMIYAHU, Zohar [IL/IL]; 30 Carcom Street, Zur Hadasa 99875 (IL).

- (74) Agent: WOLFF, BREGMAN AND GOLLER; P.O. Box 1352, Jerusalem 91013 (IL).
- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: ORGANO-CLAY COMPLEXES

(57) Abstract: The invention provides a process for the preparation of a pigment which is an organo-clay complex comprising introducing an aniline derivative acidified by hydrochloric acid and a secondary aromatic compound into a smectite mineral and treating the same with sodium nitrite solution whereby an azo-dye compound is obtained in the interlayer space of the clay to form pigmented particles.



ORGANO-CLAY COMPLEXES

The present invention relates to a process for the preparation of a pigment which is an organo-clay complex. More particularly, the present invention relates to the in-situ preparation of azo-dye compounds adsorbed onto clay particles.

Clay minerals are famous for their ability to adsorb organic compounds. Among the clay minerals, montmorillonite and other smectite minerals are the most useful for this purpose. The alumino- or magnesium silicate framework has a layer structure and in the crystal the layers are arranged one above the other. Each layer is negatively charged and the electric charge is neutralized by exchangeable cations, most of which are located in the interlayer space. The adsorbed organic cations and molecules are also located in the interlayer space.

Organic cationic dyes are adsorbed by the smectite minerals causing the mineral to be colored. The colored mineral may be used as a pigment for the color industry. The subject was recently reviewed Yariv S. (2001) Staining of clay minerals and visible adsorption spectroscopy of dye-clay complexes in "Organo-clay complexes and interactions" (S. Yariv and H. Cross, editors) Marcel Dekker, Inc., New York, pp. 463-566. The adsorption of these dyes takes place mainly by the mechanism of cation exchange.

Azo-dyes are used for different color applications. They are synthesized from aniline or its primary amine derivative, a second aromatic compound and sodium nitrite to yield products of the general formula Ar'-N=N-Ar", called azo compounds. In this reaction, known as coupling, the nitrogen of the diazonium group is retained in the product.

$$Ar'N_2 + Ar''H \rightarrow Ar'-N=N-Ar'' + H^+$$

The aromatic ring of the second compound must, in general, contain a powerfully electron-releasing group, generally —NHR or —NH₂.

There are three papers in the literature which describe the adsorption of azodyes by expanding clay minerals. Siffert, B. (1978) Preparation et etude spectrometrique de complexessilicates phyllitteux colorants azoiques. *Clay Miner.*, pp. 13: 147-165. Siffert studied vermiculite complexes of the cationic azo-dye chrysoidine and the molecular azo-dyes *para*-dimethylaminoazobenzene, bis(*para*-dimethyl-aminophenylazo)-orthotolidine, and bis(*para*-dimethyl-

2

aminophenylazo)Diphenyl. Depending on the pH of the system, the dye adsorption occurs either by molecular or by cation exchange mechanism.

Gabelica, Z., Valange, S., Shibata, M., Hotta, H. (2001): Stability against color fading of azo-dyes encapsulated in Ca-aluminosilicate mesoporous substraes. *Microporous and Mesoporous Materials*, 44-45: 645-652. Gabelica et al. studied the adsorption of Lithol Rubine B by the synthetic clay MCM-41.

Yermiyahu, Z., Lapides, I., Yariv, S. (2000). Adsorption of Congo-Red by montmorillonite. *Proc. Isr. Geological Soc., Annual Meeting, Ma'a lot 2000 (Abstracts)*, p. 131. Yermiyahu, et al. studied the adsorption of Congo-Red by montmorillonite. This azo-dye is adsorbed by the clay mineral although it is an anionic species.

According to the present invention it has now been discovered that azo-dyes can be formed in-situ within natural and synthetic clay minerals of the smectite family, e.g., montmorillonite.

Thus according to the present invention there is now provided a process for the preparation of a pigment which is an organo-clay complex comprising introducing an aniline derivative acidified by hydrochloric acid and a secondary aromatic compound into a smectite mineral and treating the same with sodium nitrate solution whereby an azo-dye compound is obtained in the interlayer space of the clay to form pigmented particles.

In preferred embodiments of the present invention said aniline derivative is selected from the group consisting of anilinium chloride, naphthyl—ammonium chloride and benzidinium chloride and said secondary aromatic compound is selected from the group consisting of 2-hydroxy-naphthalene, 2,3-dihydroxy-naphthalene, 4-amino-2-chlorotoluene-5-sulfonic acid, 2-chloro-4-aminotoluene and 2-hydroxy-1-napthoic acid.

While the invention will now be described in connection with certain preferred embodiments in the following examples so that aspects thereof may be more fully understood and appreciated, it is not intended to limit the invention to these particular embodiments. On the contrary, it is intended to cover all alternatives, modifications and equivalents as may be included within the scope of the invention as defined by the appended claims. Thus, the following examples which include preferred embodiments will serve to illustrate the practice of this invention, it being

understood that the particulars shown are by way of example and for purposes of illustrative discussion of preferred embodiments of the present invention only and are presented in the cause of providing what is believed to be the most useful and readily understood description of formulation procedures as well as of the principles and conceptual aspects of the invention.

EXAMPLES

The present pigments are prepared from different smectite minerals. Most of the work was carried out with montmorillonite and Laponite. The following examples and colors are related to montmorillonite.

Example 1

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt 1-Naphthylammonium chloride was added and stirred. After two hours 2-Chloro-4-aminotoluene was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour.

2.0 ml of aqueous solution of 0.1 molar NaNO₂ was added, and a violet color was developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

Example 2

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt 1-Naphthylammonium chloride was added and stirred. After two hours 4-Amino-2-chlorotoluene-5-sulfonic Acid was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour. 2.0 ml of aqueous solution of 0.1 molar NaNO₂ was added, and a red color was developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

Example 3

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt Anilinium-chloride was added and stirred. After two hours 2,3-Dihydroxynaphthalene was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour.

2.0 ml of aqueous solution of 0.1 molar NaNO2 was added, and a red color was

4

developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

Example 4

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt 1-Naphthylammonium chloride was added and stirred. After two hours 1-Naphthylammonium chloride was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour.

2.0 ml of aqueous solution of 0.1 molar NaNO₂ was added, and a blue color was developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

Example 5

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt 1-Naphthylammonium chloride was added and stirred. After two hours 2,3-Dihydroxynaphthalene was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour.

2.0 ml of aqueous solution of 0.1 molar NaNO₂ was added, and a violet color was developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

Example 6

To 20 ml of an aqueous 1% clay suspension, 1.0 ml of an aqueous solution of 0.1 molar of the salt Anilinium-hydrochloride was added and stirred. After two hours 2-Hydroxy-1-naphtoic acid was added (1.0 ml of an aqueous or alcoholic solution of 0.1 molar) and the mixture was stirred for half an hour.

2.0 ml of aqueous solution of 0.1 molar NaNO₂ was added, and a red color was developed immediately. The intensity of the color increased within 24 hours. After one week the solid fraction was washed three times with distilled water and was separated from the supernatant by centrifugation.

It will be evident to those skilled in the art that the invention is not limited to the details of the foregoing illustrative examples and that the present invention may be embodied in other specific forms without departing from the essential attributes

5

thereof, and it is therefore desired that the present embodiments and examples be considered in all respects as illustrative and not restrictive, reference being made to the appended claims, rather than to the foregoing description, and all changes which come within the meaning and range of equivalency of the claims are therefore intended to be embraced therein.

6

WHAT IS CLAIMED IS:

1. A process for the preparation of a pigment which is an organo-clay complex comprising introducing an aniline derivative acidified by hydrochloric acid and a secondary aromatic compound into a smectite mineral and treating the same with sodium nitrite solution whereby an azo-dye compound is obtained in the interlayer space of said clay to form pigmented particles.

- 2. A process according to claim 1 wherein said aniline derivative is selected from the group consisting of anilinium chloride, naphthyl —ammonium chloride and benzidinium chloride.
- 3. A process according to claim 1 wherein said secondary aromatic compound is selected from the group consisting of 2-hydroxy-naphthalene, 2,3-dihydroxy-naphthalene, 4-amino-2-chlorotoluene-5-sulfonic acid, 2-chloro-4-aminotoluene and 2-hydroxy-1-napthoic acid.

INTERNATIONAL SEARCH REPORT

International polication No. PCT/IL 03/00497

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C09863/00 C09841/00

C09C1/42

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols) C09B C09C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data, PAJ

Category °	Cilation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Υ	GB 665 474 A (NAT LEAD CO) 23 January 1952 (1952-01-23) page 1, line 66-82 page 2, line 64 -page 82; example 4	1-3
Y	HUDDERSMAN K ET AL: "AZO DYES ENCAPSULATED WITHIN ALUMINOSILICATE MICROPOROUS MATERIALS" JOURNAL OF THE SOCIETY OF DYERS AND COLOURISTS, SOCIETY OF DYERS AND COLOURISTS. BRADFORD, GB, vol. 114, no. 5/6, 1 May 1998 (1998-05-01), pages 155-159, XP000751702 ISSN: 0037-9859 the whole document	1-3

Further documents are listed in the continuation of box C.	χ - Patent family members are listed in annex.
Special categories of cited documents: A' document defining the general state of the art which is not considered to be of particular relevance E' earlier document but published on or after the international filing date L' document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another cliation or other special reason (as specified) O' document referring to an oral disclosure, use, exhibition or other means P' document published prior to the international filing date but later than the priority date claimed	 "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search 20 October 2003	Date of malling of the international search report . 03/11/2003
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Authorized officer Dauksch, H



International explication No PCT/IL U3/00497

		PCT/IL 03	/00497
C.(Continu	etion) DOCUMENTS CONSIDERED TO BE RELEVANT	· · · · · · · · · · · · · · · · · · ·	
Category °	Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No.
A	CHEMICAL ABSTRACTS, vol. 117, no. 2, 13 July 1992 (1992-07-13) Columbus, Ohio, US; abstract no. 8561p, KOWALSKA M. ET AL: "Interactions of montmorillonite with p-nitro- and p-methoxyanilines" XP002258497 abstract		1
•	&.CLAYS CLAY_MINER., vol. 40, no. 2, 1992, pages 237-239,		
		·	
et ·			





International explication No.

				i	PC1/1L -03/0049/		
Patent document cited in search report		Publication date		Patent family member(s)		Publication date	·
GB 665474	Α	23-01-1952	NONE				
		ان سا سا سا _ن ہیں۔ ہیں بند ابن کی کے بعد ہ					
			•				
•							
			•				
					•		